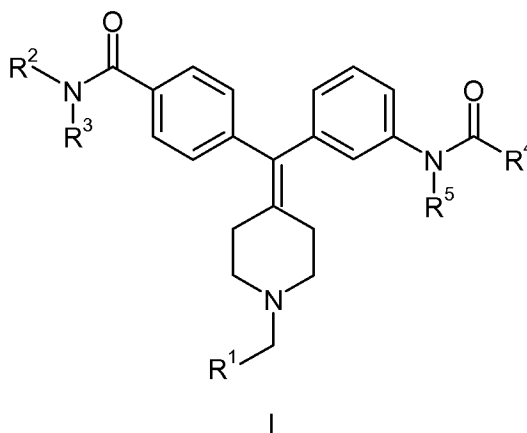


## Listing of Claims

This listing of claims will replace all prior versions, and listings, of claims in the application:

1. (currently amended) A compound of formula I, a pharmaceutically acceptable salt thereof, diastereomers, enantiomers, or mixtures thereof:



wherein

$R^1$  is ~~selected from~~  $C_{6-10}$ aryl and or  $C_{2-6}$ heteroaryl, wherein said  $C_{6-10}$ aryl and  $C_{2-6}$ heteroaryl are optionally substituted with one or more groups selected from  $C_{1-6}$ alkyl,  ~~$-R$ ,  $-NO_2$ ,  $-OR$ ,  $-O-C_{1-6}$ alkyl,  $-Cl$ ,  $-Br$ ,  $-I$ ,  $-F$ , and  $-CF_3$ ,  $C(=O)R$ ,  $C(=O)OH$ ,  $NH_2$ ,  $SH$ ,  $NHR$ ,  $NR_2$ ,  $SR$ ,  $SO_3H$ ,  $SO_2R$ ,  $S(=O)R$ ,  $CN$ ,  $OH$ ,  $C(=O)OR$ ,  $C(=O)NR_2$ ,  $NRC(=O)R$ , and  $NRC(=O)OR$~~ , wherein  $R$  is, independently, a hydrogen or  $C_{1-6}$ alkyl; and

$R^2$ ,  $R^3$ ,  $R^4$  and  $R^5$  are, independently, selected from hydrogen,  $C_{1-6}$ alkyl, and  $C_{3-6}$ cycloalkyl, wherein said  $C_{1-6}$ alkyl and  $C_{3-6}$ cycloalkyl are optionally substituted with one or more groups selected from  $C_{1-6}$ alkyl,  ~~$-R$ ,  $-NO_2$ ,  $-OR$ ,  $-O-C_{1-6}$ alkyl,  $-Cl$ ,  $-Br$ ,  $-I$ ,  $-F$ , and  $-CF_3$ ,  $C(=O)R$ ,  $C(=O)OH$ ,  $NH_2$ ,  $SH$ ,  $NHR$ ,  $NR_2$ ,  $SR$ ,  $SO_3H$ ,  $SO_2R$ ,  $S(=O)R$ ,  $CN$ ,  $OH$ ,  $C(=O)OR$ ,  $C(=O)NR_2$ ,  $NRC(=O)R$ , and  $NRC(=O)OR$~~ , wherein  $R$  is, independently, a hydrogen or  $C_{1-6}$ alkyl.

2. (currently amended) A compound according to claim 1,

wherein  $R^1$  is ~~selected from~~ phenyl; pyridyl; thienyl; furyl; imidazolyl; triazolyl; pyrrolyl; thiazolyl; and or N-oxido-pyridyl, wherein  $R^1$  is optionally substituted with one or more groups selected from  $C_{1-6}$ alkyl, ~~halogenated  $C_{1-6}$ alkyl~~,  $-NO_2$ ,  $-CF_3$ ,  $C_{1-6}$  alkoxy, chloro, fluoro, bromo, and iodo;

$R^2$ ,  $R^3$ , and  $R^4$  are, independently,  $C_{1-3}$ alkyl or halogenated  $C_{1-3}$ alkyl; and

R<sup>5</sup> is ~~selected from~~ hydrogen, C<sub>1-6</sub>alkyl, ~~and or~~ C<sub>3-6</sub>cycloalkyl, wherein said C<sub>1-6</sub>alkyl and C<sub>3-6</sub>cycloalkyl are optionally substituted with one or more groups selected from C<sub>1-6</sub>alkyl, ~~halogenated C<sub>1-6</sub>alkyl~~, -NO<sub>2</sub>, -CF<sub>3</sub>, C<sub>1-6</sub> alkoxy, chloro, fluoro, bromo, and iodo.

3. (currently amended) A compound according to claim 1,

wherein R<sup>1</sup> is ~~selected from~~ phenyl; pyridyl; thienyl; furyl; imidazolyl; pyrrolyl; ~~and or~~ thiazolyl, wherein R<sup>1</sup> is optionally substituted with one or more groups selected from C<sub>1-6</sub>alkyl, ~~halogenated C<sub>1-6</sub>alkyl~~, -NO<sub>2</sub>, -CF<sub>3</sub>, C<sub>1-6</sub> alkoxy, chloro, fluoro, bromo, and iodo;

R<sup>2</sup>, R<sup>3</sup>, and R<sup>4</sup> are, independently, C<sub>1-3</sub>alkyl or halogenated C<sub>1-3</sub>alkyl; and

R<sup>5</sup> is hydrogen.

4. (original) A compound according to claim 1,

wherein R<sup>1</sup> is selected from phenyl, pyridyl, thienyl, furyl, imidazolyl, pyrrolyl, and thiazolyl;

R<sup>2</sup> and R<sup>3</sup> are ethyl;

R<sup>4</sup> is C<sub>1-3</sub>alkyl; and

R<sup>5</sup> is hydrogen.

5. (original) A compound according to claim 1, wherein the compound is selected from:

4-[[3-(acetylamino)phenyl][1-(thien-2-ylmethyl)piperidin-4-ylidene]methyl]-N,N-diethylbenzamide;

4-[[3-(acetylamino)phenyl][1-(2-furylmethyl)piperidin-4-ylidene]methyl]-N,N-diethylbenzamide;

4-[[3-(acetylamino)phenyl][1-(phenylmethyl)-4-piperidinyldiene]methyl]-N,N-diethyl-benzamide;

4-[[3-(acetylamino)phenyl][1-(3-thienylmethyl)-4-piperidinyldiene]methyl]-N,N-diethyl-benzamide;

4-[[3-(acetylamino)phenyl][1-(3-pyridinylmethyl)-4-piperidinyldiene]methyl]-N,N-diethyl-benzamide;

4-[[3-(acetylamino)phenyl][1-(4-pyridinylmethyl)-4-piperidinyldiene]methyl]-N,N-diethyl-benzamide;

4-[[3-(acetylamino)phenyl][1-(pyridin-2-ylmethyl)piperidin-4-ylidene]methyl]-N,N-diethylbenzamide;

4-[[3-(acetylamino)phenyl][1-(1,3-thiazol-4-ylmethyl)piperidin-4-ylidene]methyl]-N,N-diethylbenzamide;

4-[[3-(acetylamino)phenyl][1-(1,3-thiazol-5-ylmethyl)piperidin-4-ylidene]methyl]-N,N-diethylbenzamide; and pharmaceutically acceptable salts thereof.

6. (cancelled)

7. (withdrawn) A method for the therapy of pain, anxiety or functional gastrointestinal disorders comprising the step of administering to said animal in need of such therapy a therapeutically effective amount of a compound according to claim 1.

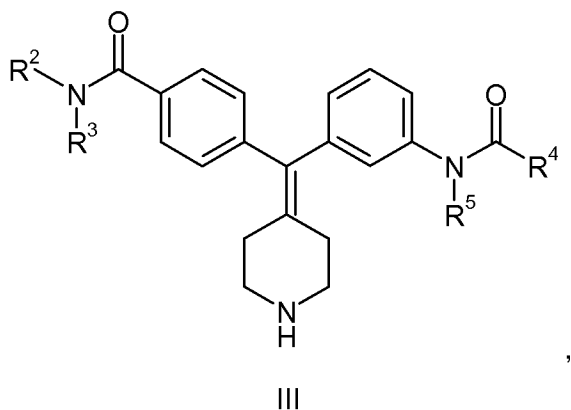
8. (previously presented) A pharmaceutical composition comprising a compound according to claim 1 and a pharmaceutically acceptable carrier.

9. (withdrawn) A method for the therapy of pain in a warm-blooded animal, comprising the step of administering to said animal in need of such therapy a therapeutically effective amount of a compound according to claim 1.

10. (withdrawn) A method for the therapy of functional gastrointestinal disorders in a warm-blooded animal, comprising the step of administering to said animal in need of such therapy a therapeutically effective amount of a compound according to claim 1.

Claims 11-12. (cancelled)

13. (original) A compound of formula III:



wherein

$R^2$ ,  $R^3$ ,  $R^4$  and  $R^5$  are, independently, selected from hydrogen,  $C_{1-6}$ alkyl, and  $C_{3-6}$ cycloalkyl, wherein said  $C_{1-6}$ alkyl and  $C_{3-6}$ cycloalkyl are optionally substituted with one or more groups selected from  $C_{1-6}$ alkyl,  $-R$ ,  $-NO_2$ ,  $-OR$ ,  $-O-C_{1-6}$ alkyl,  $-Cl$ ,  $-Br$ ,  $-I$ ,  $-F$ , and  $-CF_3$ ,  $C(=O)R$ ,  $C(=O)OH$ ,  $NH_2$ ,  $SH$ ,  $NHR$ ,  $NR_2$ ,  $SR$ ,  $SO_3H$ ,  $SO_2R$ ,  $S(=O)R$ ,  $CN$ ,  $OH$ ,  $C(=O)OR$ ,  $C(=O)NR_2$ ,  $NRC(=O)R$ , and  $NRC(=O)OR$ , wherein  $R$  is, independently, a hydrogen or  $C_{1-6}$ alkyl.

14. (withdrawn) A method for the therapy of pain, anxiety or functional gastrointestinal disorders comprising the step of administering to said animal in need of such therapy a therapeutically effective amount of a compound according to claim 2.

15. (withdrawn) A method for the therapy of pain, anxiety or functional gastrointestinal disorders comprising the step of administering to said animal in need of such therapy a therapeutically effective amount of a compound according to claim 3.

16. (withdrawn) A method for the therapy of anxiety comprising the step of administering to said animal in need of such therapy a therapeutically effective amount of a compound according to claim 1.

17. (withdrawn) A method for the therapy of anxiety comprising the step of administering to said animal in need of such therapy a therapeutically effective amount of a compound according to claim 2.

18. (withdrawn) A method for the therapy of anxiety comprising the step of administering to said animal in need of such therapy a therapeutically effective amount of a compound according to claim 3.

19. (previously presented) A pharmaceutical composition comprising a compound according to claim 2 and a pharmaceutically acceptable carrier.

20. (previously presented) A pharmaceutical composition comprising a compound according to claim 3 and a pharmaceutically acceptable carrier.

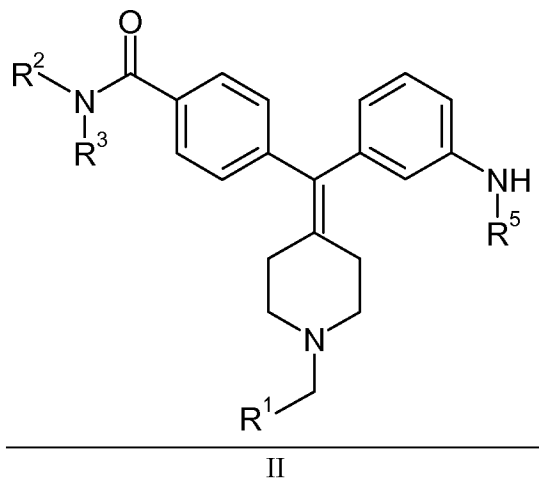
21. (previously presented) A pharmaceutical composition comprising a compound according to claim 4 and a pharmaceutically acceptable carrier.

22. (previously presented) A pharmaceutical composition comprising a compound according to claim 5 and a pharmaceutically acceptable carrier.

23. (previously presented) A compound according to claim 13, wherein the compound is 4-[[3-(acetylamino)phenyl](piperidin-4-ylidene)methyl]-N,N-diethylbenzamide.

24. (new) A process for preparing a compound of formula I according to claim 1, comprising:

reacting a compound of formula II with  $X-C(=O)-R^4$  or  $R^4C(=O)-OC(=O)R^4$ :



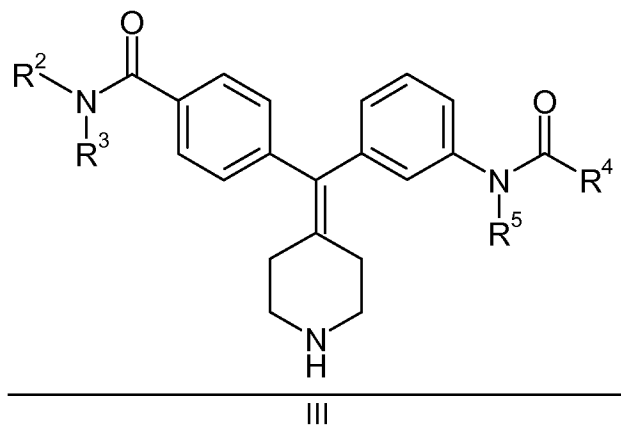
wherein

$R^1$  is  $C_{6-10}$ aryl or  $C_{2-6}$ heteroaryl, wherein said  $C_{6-10}$ aryl and  $C_{2-6}$ heteroaryl are optionally substituted with one or more groups selected from  $C_{1-6}$ alkyl,  $-O-C_{1-6}$ alkyl,  $-Cl$ ,  $-Br$ ,  $-I$ ,  $-F$ , and  $-CF_3$ ;

$R^2$ ,  $R^3$ ,  $R^4$  and  $R^5$  are, independently, selected from hydrogen,  $C_{1-6}$ alkyl, and  $C_{3-6}$ cycloalkyl, wherein said  $C_{1-6}$ alkyl and  $C_{3-6}$ cycloalkyl are optionally substituted with one or more groups selected from  $C_{1-6}$ alkyl,  $NO_2$ ,  $-O-C_{1-6}$ alkyl,  $-Cl$ ,  $-Br$ ,  $-I$ ,  $-F$ , and  $-CF_3$ ; and

$X$  is  $Cl$ ,  $Br$  or  $I$ .

25. (new) A process for preparing a compound of formula I, according to claim 1 comprising:  
reacting a compound of formula III with  $R^1-CHO$  or  $R^1-CH_2X$ :



wherein

$R^1$  is  $C_{6-10}$ aryl or  $C_{2-6}$ heteroaryl, wherein said  $C_{6-10}$ aryl and  $C_{2-6}$ heteroaryl are optionally substituted with one or more groups selected from  $-R$ ,  $-NO_2$ ,  $-OR$ ,  $-Cl$ ,  $-Br$ ,  $-I$ ,  $-F$ ,  $-CF_3$ ,  $-C(=O)R$ ,  $-C(=O)OH$ ,  $-NH_2$ ,  $-SH$ ,  $-NHR$ ,  $-NR_2$ ,  $-SR$ ,  $-SO_3H$ ,  $-SO_2R$ ,  $-S(=O)R$ ,  $-CN$ ,  $-OH$ ,  $-C(=O)OR$ ,  $-$

C(=O)NR<sub>2</sub>, -NRC(=O)R, and -NRC(=O)-OR, wherein R is, independently, a hydrogen or C<sub>1-6</sub>alkyl;

R<sup>2</sup>, R<sup>3</sup>, R<sup>4</sup> and R<sup>5</sup> are, independently, selected from hydrogen, C<sub>1-6</sub>alkyl, and C<sub>3-6</sub>cycloalkyl, wherein said C<sub>1-6</sub>alkyl and C<sub>3-6</sub>cycloalkyl are optionally substituted with one or more groups selected from C<sub>1-6</sub>alkyl, -NO<sub>2</sub>, -O-C<sub>1-6</sub>alkyl, -Cl, -Br, -I, -F, and -CF<sub>3</sub>; and

X is Cl, Br or I.